IN THE CLAIMS

1-42. (Canceled)	
43.	(Currently Amended) A pharmaceutical composition comprising:
	a therapeutically effective amount of an antigen presenting cell pulsed with an inactivated
non-recombinant human immunodeficiency virus (HIV); and	
	a pharmaceutically acceptable carrier;
	wherein the inactivated human immunodeficiency virus is chemically inactivated by 2,2'-
dithiopyridine, and wherein the composition expands in vivo expression of virus-specific CD8+T	
cells, a	and said virus-specific CD8+ cells kill HIV-infected cells; and

44. (Currently Amended) A pharmaceutical composition comprising:

plastic-adherence followed by culture with GM-CSF and IL-4.

Please replace the previous version of the claims with the following:

a therapeutically effective amount of an antigen presenting cell pulsed with an inactivated non-recombinant human immunodeficiency virus (HIV); and

wherein said antigen presenting cell is a dendritic cell obtained from a monocyte by

a pharmaceutically acceptable carrier;

wherein the inactivated human immunodeficiency virus is chemically inactivated by 2,2'-dithiopyridine, and wherein the composition expands *in vivo* expression of virus-specific CD8+T cells, and said virus-specific CD8+ cells kill HIV-infected cells,

wherein said antigen presenting cell is a dendritic cell obtained from a monocyte by plastic-adherence followed by culture with GM-CSF and IL-4, and

— wherein said inactivated human immunodeficiency virus is an inactivated autologous human immunodeficiency virus.

- 45. (Cancelled)
- 46. (Currently Amended) The pharmaceutical composition of Claim [[45]] 43, wherein said dendritic cell is an autologous dendritic cell.
- 47-51. (Cancelled)
- 52. (Previously Presented) The pharmaceutical composition of Claim 43, further comprising an adjuvant for optimizing the virus-specific CTL response.
- 53. (Previously Presented) The pharmaceutical composition of claim 52, wherein said adjuvant is a protease inhibitor.
- 54. (Previously Presented) The pharmaceutical composition of claim 53, wherein said protease inhibitor is indinavir.
- 55. (Previously Presented) The pharmaceutical composition of claim 54, wherein said composition comprises non-antiviral concentration of indinavir.
- 56. (Previously Presented) The pharmaceutical composition of claim 55, comprising a concentration of indinavir of 10 nM.